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(54) Ultraviolet radiation-curable coatings for optical fibers and optical fibers coated therewith

UV-Strahlungshärtbare Beschichtung für optische Fasern und damit beschichtete optische Fasern

Revêtements durcissables par irradiation ultraviolette pour fibres optiques et fibres optiques revêtus
par ceux-ci

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Description**BACKGROUND OF THE INVENTION:**

5 The present invention relates to radiation-curable primary coatings for optical fibers and to optical fibers containing such coatings.

Until recently, the optical fiber industry was concentrated on so-called "long haul" applications, wherein optical fibers were used to traverse long distances such as in transoceanic or transcontinental cables. In such applications, optical fibers required shielding with voluminous protective cabling material in sheltered subterranean or submarine environments and thus were not directly exposed to environmental hazards.

10 A recent trend in the optical fiber market is in local area networks for fiber-to-the-home uses. The fibers in such uses are directly exposed to much harsher conditions than previous applications of glass fibers, including severe temperature and humidity extremes. Consequently, previously used coatings did not perform well under such adverse conditions; hence, a need existed for the development of higher performance coatings. Such coatings needed to be able to withstand the above conditions, i.e., to possess thermal, oxidative and hydrolytic stability, and to protect the fiber over 15 the long term, i.e., over twenty-five years' time.

15 Optical fibers used for light transmission can be prepared which, immediately after drawing, are exceptionally strong and have very few intrinsic defects. However, such pristine fibers are very easily flawed by exposure to environmental conditions including dust and moisture. Even a small flaw can reduce the strength of a fiber by an order of magnitude, rendering it brittle and easily broken by a weak external forces. Therefore, optical glass fibers have, in the prior art, been clad with at least one resin coat immediately after their preparation, whose minimum requirement is to protect 20 the underlying pristine fiber from such external forces.

25 Typically, at least two coatings, a primary, or buffer, coating and a secondary coating, have been used. The inner, or primary, coating is applied directly to the glass fiber and, when cured, forms a soft, rubbery, compliant material which serves as a buffer to cushion and protect the fiber by relieving the stresses created when the fiber is bent, cabled or spooled. Such stress might otherwise induce microbending of the fibers and cause attenuation of the light traveling through them, resulting in inefficient signal transmission. The secondary coating is applied over the primary coating, and must function as a hard, tough protective outer layer preventing damage to the glass fiber during processing and use.

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PRIMARY COATING

Certain characteristics are desirable for the primary coating, and others for the secondary coating. For example, the primary coating must maintain adequate adhesion to the glass fiber during thermal and hydrolytic aging, yet be 35 strippable therefrom for splicing purposes. The tensile modulus of the primary coating must be low to cushion and protect the fiber by readily relieving the stresses on the fiber which can induce microbending and consequent inefficient signal transmission. This cushioning effect must be maintained through the temperature range in which the fibers may be exposed throughout their lifetime. Thus, it is necessary for the primary coating to have a low glass transition temperature (T_g). This low glass transition temperature will ensure that the coating remains in its rubbery state throughout the 40 possible use temperature range.

The primary coating should also have a relatively high refractive index, i.e., greater than that of the cladding material of the fiber to be coated. This high refractive index allows for a refractive index differential between the glass cladding and the primary coating. This differential allows errant light signals to be refracted away from the glass core.

Another requisite quality of the primary (buffer) coating is resistance to moisture. Moisture will rapidly degrade the 45 strength of the coating itself as well as the underlying glass fibers under stress. The reaction is one of hydrolysis and stress corrosion. Moisture will also adversely affect the adhesion of the primary (buffer) coating to the glass, resulting in possible delamination. It is therefore desirable for the coating to be as hydrophobic as possible. Preferably, the primary coating should have a water absorption value of less than 5% by weight, and more preferably less than 2.5% by weight.

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SECONDARY COATING

Similarly, the secondary coating must have a number of essential and optimal qualities. The secondary coating must function as a hard protective layer which allows the fiber not to be damaged during its processing and use. Consequently, this secondary coating should have a relatively high glass transition temperature (T_g), i.e., about 50°C, and a high tensile modulus, i.e., about 689 MPa (100,000 psi) at 25°C. It is desirable for the secondary coating to have a T_g 55 higher than its highest use temperature, because at or near the T_g of a polymer, many physical properties such as modulus, tensile strength, thermal expansion coefficient, moisture absorptivity, and so forth, change dramatically with small

changes in temperature. This results in large changes in the fiber characteristics.

Furthermore, like the primary coating, the secondary coating should undergo minimal changes in physical properties on exposure to moisture. Many polymeric coating materials experience significant hydrolysis, plasticization, softening, and loss of protective function in the presence of water. Therefore, it is desirable that the secondary coating should also be as hydrophobic as possible. A hydrophobic secondary coating also enhances the ability of the primary coating to keep water away from the glass fiber itself, and thus inhibits moisture induced hydrolysis and stress corrosion of the glass fiber.

Also, the secondary coating must have a high refractive index, i.e., preferably higher than that of the cladding material of the glass fiber, to enable it to strip out errant light, as with the primary coating, especially in embodiments where the primary coating is omitted.

Another important property of secondary coatings is the coefficient of friction (COF). A low COF facilitates winding and unwinding of the fibers on spools. It also allows the fibers to slide easily along each other in a cable structure, thus relieving stress. However, if the COF is too low, the fibers will have a reduced tendency to stay aligned on the spool. The COF is largely determined by the cure properties of the secondary coating; if the coating has a good hard surface cure, its COF will tend to be low. Various slip additives can be added to lower the COF; however, deleterious effects such as surface tension problems, contamination, volatilization and blooming may result from their use.

DESIRABLE FEATURES FOR BOTH COATINGS

Still other desirable properties exist which apply to both the primary and secondary coatings. For example, fiber manufacturers are motivated to coat the fibers as rapidly as possible to attain the economy of faster cure speeds, as these result in higher line speeds. The cure speeds of coating materials may be determined by constructing a UV dose versus modulus curve. The lowest UV dose at which the coating modulus exhibits dose independence is considered its cure speed. There is therefore a demand for faster curing coatings; for example, high line speeds are obtained with primary and secondary coatings which may be applied wet-on-wet and simultaneously ultraviolet light-cured. One way of doing this is described in U.S. Patent 4,474,830, issued October 2, 1984 to Carl R. Taylor, which patent is expressly incorporated herein by reference.

According to this patent, an optical fiber is passed through a coating applicator comprising first and second dies. The first die confines a first coating liquid over a portion of the fiber's length. A second coating liquid is applied onto the fiber through a clearance between the first and second dies. The clearance is sufficiently small so that substantially no recirculation of the second coating liquid occurs in the vicinity of the point of application to the fiber. The second coating liquid is preferably applied at a free surface; that is, it is not in contact with either the first or second dies in the immediate vicinity of the point of application to the fiber. Additional coatings can be applied in a similar manner.

Another desirable objective for both primary and secondary UV-curable coatings is to minimize the amount of unbound material in the coating after cure. Even when the cured coatings are considered 100% solids, there may still exist a small amount of material which does not chemically bind into the polymer network on curing. Examples of such materials used in the coatings include unreacted monomer, unreacted photoinitiator, certain non-functional additives and so forth. The presence of excessive amounts of such materials is undesirable, inasmuch as volatilization of such components over time may change the physical properties of the coating. For example, volatile materials from the primary coating may permeate into the secondary coating, tending to plasticize it and resulting in strength loss. Also, volatile materials may cause production of unpleasant odors.

Still other important qualities of both optical fiber coatings are viscosity and shelf life. Good shelf life is considered formulation stability of at least six to twelve months. Viscosity can typically be somewhat adjusted by regulation of the temperature at which the coatings are applied. However, it is advantageous to set the viscosity high enough so as to maintain proper rheology and handling of the coating on application, but low enough to facilitate bubble release and to minimize the amount of heat needed in the preparation. Excessive heating is undesirable inasmuch as it may result in premature gelling or viscosity increase due to possible thermal initiation of polymerization.

PRIOR ART COATINGS

Various single or double layer fiber coatings exist in the prior art. Among these are epoxy- or urethane-based resins. However, many of these resins cure slowly; have poor hydrophobicity or poor hydrolytic, thermal and oxidative stability; and have undesirable yellowing properties.

There have also been developed primary (buffer) coatings which cure on exposure to ultraviolet radiation. Such prior art primary coatings, however, have conventionally not been very moisture resistant and have some of the same deficiencies as above.

To obviate these flaws, the primary-coated optical fibers of the prior art have been topcoated with a tough and flexible overcoat which possesses superior resistance to moisture abrasion. Prior art coatings have included extruded

nylon "jacket" coatings, which are, however, more expensive and more difficult to apply than would be an ultraviolet-cured coating.

Therefore, the present invention seeks to provide primary coatings which, when cured, have maximal thermal, oxidative and hydrolytic stability, wherein the primary coating is adequately adherent to the glass fiber yet strippable therefrom; provides adequate cushioning of the fiber; has a relatively low glass transition temperature; and is capable of relieving stress upon the fiber.

The coating can be formulated for wet-on-wet application, with simultaneous radiation curing at commercially acceptable cure speeds.

In U.S. Serial No. 742,531, filed August 8, 1991, which was a continuation of U.S. Serial No. 350,239, filed May 11, 10 1989, which is the parent application of this application, coating compositions prepared from acrylate- or methacrylate-based components and having excellent properties vis-à-vis the above characteristics were disclosed. That application discloses acrylate-modified primary and secondary coatings, the primary coating including an acrylated urethane oligomer; an alkyl acrylate monomer; a refractive index modifier; an adhesion promoter; and a photoinitiator. The secondary coating includes an acrylated urethane oligomer based on a polyester; a (meth)acrylated compound; and a photoinitiator.

WO 90/13523 discloses a liquid, radiation-cured coating composition for producing coatings on glass surfaces which contains 56 to 89% by weight of at least one diethylenic-unsaturated polyurethane, 3 to 30% by weight of at least one ethylenic-unsaturated monomer, 0.5 to 8% by weight of at least one photoinitiator and 0.05 to 6% by weight of an alkoxy silane.

WO 92/04388 teaches optical fibre coatings prepared from compositions containing a vinyl ether oligomer prepared by reacting a hydroxyl-terminated polyester or polyether, a diisocyanate and a hydroxy monovinyl ether, with mono or multifunctional vinyl ether terminated monomers, which may be derived from esters or alcohols.

EP-A-0336653 describes optical fibre coatings with a low glass transition temperature which are prepared from acrylated urethane oligomers having a molecular weight of 2,000 to 6,000, an aliphatic monofunctional acrylate or mono-functional aryl-containing acrylate having glass transition temperatures below -20°C, a photoinitiator and, optionally, a crosslinking acrylate ester having at least two acrylate or methacrylate groups per molecule and having a molecular weight less than 4,000.

SUMMARY OF THE INVENTION

It has now been discovered that it is the functional backbones of these components, rather than their end groups, that confer the excellent properties upon the compositions of the parent application. Thus, systems analogous to the acrylate- or methacrylate-based compositions of the parent application, but bearing any reactive end groups, are equally suitable in meeting these requirements. These applications are expressly incorporated by reference herein.

The present invention therefore addresses analogous systems incorporating various other exemplary end groups capable of reacting upon irradiation or other means, either by free radical initiation or cationic cure, to provide excellent performance coatings. Such end groups include, but are by no means limited to, free radical systems such as thiolene systems (based on the reaction of multifunctional thiols and unsaturated polyenes, such as vinyl ethers; vinyl sulfides; allylic ethers; and bicyclic enes); amine-ene systems (based on the reaction of multifunctional amines and unsaturated polyenes); acetylenic systems; systems wherein the reactive portion of the component is internal rather than terminal; other vinylic (e.g., styrenic) systems; acrylamide systems; allylic systems; itaconate systems and crotonate systems; and cationic cure systems such as onium salt-induced vinyl ether systems and epoxy-terminated systems which react by ring-opening; and any others based on compounds possessing reactive termini. In fact, virtually any end groups which cure by irradiation or other means but do not adversely effect the desirable properties (i.e., the oxidative, thermal and hydrolytic stability and the moisture resistance) of the cured composition are envisioned.

Another contribution of the present invention over that of the parent application is the discovery that two of the components said in the parent application to be required are, in some circumstances, optional.

One of these components is a silane adhesion promoter, which, it has now been discovered, is not required in moderate- or low-temperature, low-humidity environments.

Also, a photoinitiator is, in some instances, optional; for instance, when electron beam cure of a free radical system is envisioned, a photoinitiator may be omitted.

Accordingly, in preferred embodiments of the invention, a broad spectrum of radiation-curable primary coatings for optical fibers has been provided. The primary coatings have refractive indices greater than those of their fiber substrates, and relatively low glass transition temperatures and tensile moduli (i.e., less than about -20°C and less than about 3.44 MPa (500 psi) at 25°C, respectively). An optical fiber coated with the primary coatings of the present invention, has also been developed.

The radiation-curable primary coating for an optical fiber of the present invention comprises :

(A) from about 10 percent to about 90 percent by weight of a reactively terminated urethane oligomer which is the reaction product of (i) a hydrocarbon polyol, the hydrocarbon portion of which is from about 500 to about 4,000 molecular weight; (ii) an aliphatic polyisocyanate; and (iii) an endcapping monomer capable of supplying a reactive terminus;

(B) from about 10 percent to about 90 percent by weight of a hydrocarbon monomer containing between 6 and 20 carbon atoms which is terminated with at least one end group capable of reacting with the reactive terminus of (A);

(C) optionally, from about 0.05 percent to about 10.0 percent by weight of a photoinitiator; and

(G) from 5% to 60% by weight of a monomer or oligomer selected from the group consisting of polyalkylene glycol nonylphenyl ether acrylates, polyalkylene glycol nonylphenyl ether methacrylates, and mixtures thereof,

all of the above stated percentages by weight for the primary coating being based on the total weight of (A), (B) and (C).

Preferably, the hydrocarbon polyol portion of the oligomer (A) has a molecular weight in the range of about 600 to about 4,000.

The primary coating may optionally contain other materials, such as: a crosslinking agent; an organofunctional silane adhesion promoter; a chain transfer agent, preferably a mercapto-functional chain transfer agent; and at least one stabilizer.

A coated optical fiber and process for preparing a coated optical fiber have been developed. In one embodiment, the process comprises sequentially applying to an optical fiber a primary coating layer as described above, a secondary coating layer and radiation-curing the coating layers.

The coating compositions of the invention are also useful for other coating and protective purposes. They can be formulated to be useful on glass, ceramic, granite, and marble surfaces, and the like.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention relates in part to radiation-curable primary coatings for optical fibers. The optical fibers which are coated may comprise a glass core and a glass cladding layer. The core, for example, may comprise silica doped with oxides of germanium or phosphorus and the cladding, a pure or doped silicate such as fluorosilicate. Alternatively, the fibers may comprise a polymer-clad silica glass core. Examples of such polymer claddings include organosiloxanes such as polydimethylsiloxane or a fluorinated acrylic polymer.

The primary coating should have, *inter alia*, the following properties when cured: moisture resistance; ease of application and stripping; low volatiles content; low tensile modulus over the life of the fiber; low glass transition temperature; and long shelf life.

The coating should be transparent; nonmalodorous; fast curing; and remain adherent, even upon aging in high heat and humidity environments.

The present invention involves the recognition that it is the backbone portions of the essential (and optional) ingredients of the composition that confer the excellent performance properties upon the composition, and that any end groups meeting certain requirements may be used in formulating the various components of the present invention. It is merely required that the components contain end groups which react upon exposure to irradiation or other initiation, i.e., where the components contain at least one terminus or group that is capable of reacting with a terminus or group of other components. The reactive groups, when cured, must produce a composition which is stable with respect to thermal, hydrolytic and oxidative stresses to which the cured composition may be subjected.

Additionally, while the end groups must react to cure well upon exposure to irradiation or other initiation, they should not be so reactive as to cure prematurely, so that, if desired, a "one package" system would be unfeasible.

While not desiring to be limitative, two broad classes of systems which meet these criterion are discussed and exemplified herein. One such system is a free radical initiated system, many examples of which are well known in the art and several types of which will be exemplified below.

A second system is a cationic cure-type system, wherein reactive acids are generated in the presence of ultraviolet light. Such cationic curing systems are known in the art and are generally disclosed, for example, in U.S. Patent 3,981,987 and 4,981,881 to J. Crivello, assigned to General Electric Corporation and herein expressly incorporated by reference.

Again, however, these systems represent only nonlimitative examples of chemistries which are suitable for use in the present invention; any system which does not compromise the qualities of the cured composition may be used. Furthermore, "hybrid" compositions containing mixtures of different reactive systems may be used, so long as their mixture does not interfere with the cure of either system, and their mixture does not adversely effect the cured composition.

Examples of free radical systems include, but are not limited to, acrylate- and methacrylate-based systems; thio-

lene-based systems; all-allylic systems; other vinylic (e.g., styrenic) norbornenyl systems; systems; acrylamide systems; itaconate systems; crotonate systems; amine-ene systems; acetylenic systems; and internally reactive systems.

One preferred free radical system other than (meth)acrylate which is exemplified herein is a thiolene system, which involves the reaction of a mercaptan (thiol) with a polyene, wherein the two components are reacted in stoichiometric proportion.

The polyene component of the thiolene system may, theoretically, be any component containing a reactive, unsaturated group, most desirably attached to an electron-rich atom or group. Thus, a preferred polyene is a polyfunctional vinyl ether; other suitable groups include but are not limited to allylic ethers; vinyl sulfides; styrenes; acrylamides and acetylenes.

Another suitable class of compound includes bicyclic enes such as norbornene carboxylate (the reaction product of cyclopentadiene and acrylate), although their ester functionality may compromise hydrolytic stability.

With respect to the thiol (mercaptop) component, primary thiols containing 6 to 18 carbon atoms, and preferably 8 to 16 carbon atoms, such as 1,10-decane dithiol, or any linear, cyclic or branched hydrocarbon thiol containing from one to four thiol groups may be used. Primary thiols are preferred, inasmuch as they are most reactive, followed by secondary, and, finally, the least reactive tertiary thiols.

Alternatively, even higher reactivity is achieved when the sulfur atom of the mercaptan is in the vicinity of a carbonyl group, e.g., mercaptoacetate or mercaptopropionate, although hydrolytic stability may be impaired, due to the presence of the ester group.

As mentioned, *supra*, an alternative system is a cationic curing system. Such systems require an onium salt, e.g., 20 iodonium, sulfonium, arsonium, azonium, bromonium, or selenonium, which salt generates cations on exposure to ultraviolet light. Such onium salts are desirably modified by bonding thereto saturated hydrocarbon moieties, to improve their compatibility with the very hydrophobic components of the composition. Nonlimitative examples of such cationic curing reactive groups include vinyl ethers (wherein all reactive components are vinyl ether terminated); and epoxy terminated systems which react by ring opening (e.g., between cyclohexene oxide and oxirane or between glycidyl esters).

THE PRIMARY (BUFFER) COATING

The primary coating composition of the present invention contains four basic ingredients, (A), (B), (C) and (G).

A. The Hydrocarbon-Based Oligomer

The first ingredient is a specific hydrocarbon-based oligomer (A). This component comprises from about 10 percent to about 90 percent by weight of the composition base on the total weight of the (A), (B) and (C) ingredients of the composition. Preferably, this oligomer comprises from about 15 percent to about 85 percent, and more preferably about 20 percent to about 80 percent by weight of the composition, based on total weight of the (A) through (C) ingredients.

The particular hydrocarbon-based oligomer used in the present invention was chosen to impart good thermal, oxidative and hydrolytic stability to the cured system.

It has been known in the art that various types of UV-curable oligomers exist which may yield a soft, compliant, low glass transition temperature-type coating. Acrylate- or methacrylate-terminated monomers are particularly commonly used due to their ease of cure upon ultraviolet radiation. One system known in the art is acrylate-endcapped polybutadiene-type rubber or rubber-modified acrylated monomers as base resins. While these systems have excellent low temperature properties and are suitably hydrophobic, their internal carbon-carbon double bonds (unsaturation) make them susceptible to oxidation over a long period of time.

It is also known in the art to employ acrylated silicones as base resins in such compositions. While these have good low temperature properties and hydrophobicity, they are difficult to formulate with a suitably high refractive index, tend to have poor thermal stability, and may be susceptible to hydrogen outgassing which can lead to signal attenuation in fibers so coated.

Yet another system known in the art involves the use of acrylated fluorocarbons. While these are suitably hydrophobic and thermally stable, they are typically incompatible with most non-halogenated organic compounds. Additionally, they are very expensive relative to other systems.

To overcome many of the disadvantages of the prior art systems, the primary coating of the present invention utilizes a urethane system based on an hydrogenated polybutadiene backbone. In general, urethane acrylate systems based on polyethers or polyesters were usually characterized by poor water resistance and by thermal instability. Additionally, known urethane oligomers based on aromatic isocyanates displayed thermal instability and tended to yellow. While polyether-based urethane acrylates have excellent low T_g properties, they are, when used alone, generally not hydrophobic enough for optical fiber applications and are susceptible to oxidation. Polyester-based urethane acrylates have good thermal stability but are susceptible to hydrolysis. Furthermore, it has now been found that other end groups besides acrylate and methacrylate which are capable of supplying a reactive terminus to the reactants are also very

suitable for the purpose of this invention.

Thus, the primary coating of the present invention uses an oligomer which, in combination with the other components of the composition, obviates many of the above problems.

The oligomer (A) utilized in the primary coating of the present invention is the reaction product of (i) a hydrocarbon polyol, the hydrocarbon portion of which is from about 500 to about 4,000 molecular weight; (ii) an aliphatic polyisocyanate; and (iii) an endcapping monomer capable of supplying a reactive terminus.

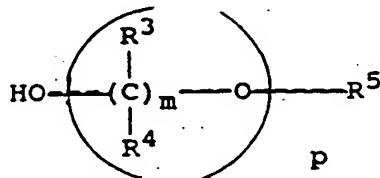
The hydrocarbon polyol (i) is provided by a linear or branched hydrocarbon containing a plurality of hydroxyl end groups, and providing a hydrocarbon backbone to the oligomer. The hydrocarbon portion is from about 500 to about 4,000 molecular weight, and preferably about 600 to about 4,000 molecular weight. Molecular weight in this case is determined by gel permeation chromatography (GPC), using a methylene chloride solvent, as measured against polystyrene molecular weight standards. By "hydrocarbon" is meant a non-aromatic compound containing a majority of methylene groups (-CH₂-) and which may contain internal unsaturation and/or pendant unsaturation. Fully saturated (i.e., hydrogenated) hydrocarbons are preferred because the long term stability of the cured optical fiber coating increases as the degree of unsaturation decreases. Suitable hydrocarbon polyols include but are not limited to hydroxyl-terminated, fully or partially hydrogenated 1,2-polybutadiene polyol; 1,2-polybutadiene polyol hydrogenated to an iodine number of from 9 to 21; fully or partially hydrogenated polyisobutylene polyol; polybutene polyol; hydrogenated dimer diols; mixtures thereof, and the like. Preferably, the hydrocarbon polyol is substantially fully hydrogenated, and thus a preferred polyol is hydrogenated 1,2-polybutadiene.

The polyisocyanate component (ii) is non-aromatic. Oligomers based on aromatic polyisocyanates effect yellowing in the cured coating. Non-aromatic polyisocyanates of from 4 to 20 carbon atoms may be employed. Suitable saturated aliphatic polyisocyanates include but are not limited to isophorone diisocyanate; dicyclohexylmethane-4,4'-diisocyanate; 1,4-tetramethylene diisocyanate; 1,5-pentamethylene diisocyanate; 1,6-hexamethylene diisocyanate; 1,7-heptamethylene diisocyanate; 1,8-octamethylene diisocyanate; 1,9-nonamethylene diisocyanate; 1,10-decamethylene diisocyanate; 2,2,4-trimethyl-1,5-pentamethylene diisocyanate; 2,2'-dimethyl-1,5-pentamethylene diisocyanate; 3-methoxy-1,6-hexamethylene diisocyanate; 3-butoxy-1,6-hexamethylene diisocyanate; omega, omega'-dipropylether diisocyanate; 1,4-cyclohexyl diisocyanate; 1,3-cyclohexyl diisocyanate; trimethylhexamethylene diisocyanate; and mixtures thereof. Isophorone diisocyanate is a preferred aliphatic polyisocyanate.

The reaction rate between the hydroxyl-terminated hydrocarbon and the diisocyanate may be increased by use of a catalyst in the amount of 100 to 200 ppm. Suitable catalysts include but are not limited to dibutyl tin dilaurate, dibutyl tin oxide, dibutyl tin di-2-hexoate, stannous oleate, stannous octoate, lead octoate, ferrous acetoacetate, and amines such as triethylamine, diethylmethylamine, triethylenediamine, dimethyl-ethylamine, morpholine, N-ethyl morpholine, piperazine, N,N-dimethyl benzylamine, N,N-dimethyl laurylamine, and mixtures thereof. A preferred catalyst is dibutyl tin dilaurate.

The endcapping monomer (iii) is one which is capable of supplying at least one reactive terminus.

An exemplary end group conforms to the formula



wherein R⁵ is selected from the group consisting of acrylic, methacrylic, vinylic, allylic, styrenic, acrylamide, norbornenyl, acetylenic, epoxy, mercapto, amino, itaconic and crotonic; R³ and R⁴ are independently selected from the group consisting of hydrogen, methyl, ethyl or propyl; m is an integer from 1 to 10, and p is 0 or 1.

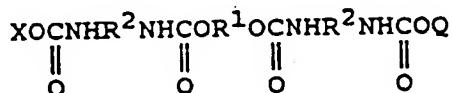
Examples of compounds which may be used for the endcapping monomer include but are not limited to acrylates,

methacrylates, vinyl ethers, vinyl sulfides, allyls, bicyclic enes, mercaptans, acetylenes, epoxides, amines, styrenes, acrylamides, and so forth. Suitable hydroxyl-terminated compounds which may be used as the endcapping monomer include but are not limited to hydroxyethyl acrylate; hydroxyethyl methacrylate; hydroxypropyl acrylate; hydroxypropyl methacrylate; hydroxybutyl acrylate; hydroxybutyl methacrylate; allyl ether; hydroxyethyl vinyl ether; hydroxypropyl vinyl ether; hydroxybutyl vinyl ether; hydroxyethyl mercaptan; hydroxypropyl mercaptan; hydroxyethyl-3-mercaptopropionate; and hydroxypropyl-3-mercaptopropionate.

In an acrylate based system, a preferred endcapping monomer is hydroxyethyl acrylate or hydroxyethyl methacrylate. In a thiolene or in a cationic cure system, a preferred end-capping monomer is hydroxybutyl vinyl ether or, in the thiolene system, hydroxypropyl mercaptan.

The molar ratio of the polyol, diisocyanate and endcapping monomer is preferably approximately 1:2:2.

A preferred oligomer is based on the formula



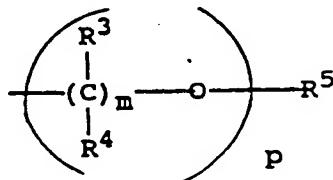
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where

R¹ is a linear or branched hydrocarbon polymer of from 500 to 4,000 (preferably 600 to 4,000) molecular weight selected from the group consisting of fully or partially hydrogenated 1,2-polybutadiene; 1,2-polybutadiene hydrogenated to an iodine number of from 9 to 21; fully or partially hydrogenated polyisobutylene; polybutene; and hydrogenated dimers;

15 R² is a linear, branched or cyclic alkylene of from six to twenty carbon atoms; and X and Q are independently either

20 (a) a radical of the formula:



30 wherein R⁵, m and p have the same meaning as above, or

(b) a saturated alkyl radical of from nine to twenty carbon atoms, with the proviso that the above oligomer must possess at least one reactive terminal (end) group.

An especially preferred monomer is one in which R⁵ is acrylic or methacrylic; this oligomer is disclosed in U.S. Patent 4,786,586 to Lee et al., which patent is herein incorporated by reference.

35 The above-described oligomer is particularly suitable because its saturated hydrocarbon backbone gives the oligomer suitable hydrophobicity and thermal, oxidative and hydrolytic stability. The glass transition temperature of the oligomer is approximately -20°C and may be lowered by blending with additives, discussed *infra*.

The primary coating containing this oligomer has a water absorption value of less than 5% by weight, and preferably less than 2.5%.

40 Another particularly preferred oligomer is the vinyl ether analog or mercapto analog of the above oligomer, i.e., where R⁵ is either vinylic or thiol, respectively. This oligomer also confers excellent thermal, oxidative and hydrolytic stability and hydrophobicity to a cured coating which includes it.

45 **B. The Hydrocarbon Monomer**

The second essential component of the composition of the primary coating composition is a hydrocarbon monomer which is terminated with at least one end group capable of reacting with the reactive terminus of (A). The hydrocarbon portion (moiety) of the monomer has between 6 and 20 carbon atoms, and preferably between 8 and 18. This monomer 50 (B) may be either straight chained, branched or cyclic. This component comprises from about 10 percent to about 90 percent by weight of the composition, based upon the total weight of the (A), (B) and (C) ingredients of the composition. Preferably, it comprises from about 15 percent to about 85 percent, and more preferably from about 20 percent to about 80 percent by weight of the composition, based upon the total weight of the components (A), (B), and (C).

This second component is one having one or more end groups capable of reacting with the reactive terminus of (A). 55 Thus, for example, in an acrylate system, the component might be acrylate-or methacrylate-terminated; in a thiolene system, vinyl ether-or mercaptan-terminated; and in a cationic cure system, vinyl ether-terminated.

The monomer is selected to be one that is compatible with the very hydrophobic oligomer discussed above. It is also chosen to be one which is soft-curing and which has a low Tg, thus lowering the Tg of the composition including

it. Suitable examples of such monomers include but are not limited to acrylates and methacrylates such as hexyl acrylate; hexyl methacrylate; 2-ethylhexyl acrylate; 2-ethylhexyl methacrylate; isoctyl acrylate; isoctyl methacrylate; octyl acrylate; octyl methacrylate; decyl acrylate; decyl methacrylate; isodecyl acrylate; isodecyl methacrylate; lauryl acrylate; lauryl methacrylate; tridecyl acrylate; tridecyl methacrylate; palmitic acrylate; palmitic methacrylate; stearyl acrylate; stearyl methacrylate; C₁₄ to C₁₅ hydrocarbon diol diacrylates; C₁₄ to C₁₅ hydrocarbon diol dimethacrylates; primary C₆ to C₁₈ hydrocarbon-based dithiols such as 1,10-decane dithiol; secondary or tertiary dithiols (which are less reactive than the primary thiols); C₆ to C₁₈ thiolated esters such as mercaptoacetate esters or mercaptopropionate esters (which are desirably reactive but which may lack in long-term hydrolytic stability); C₄ to C₁₈ hydrocarbon diol divinyl ethers such as hexanediol divinylether; lauryl vinyl ether; ethylhexyl vinyl ether; isodecyl vinyl ether; butanediol divinyl ether; cyclohexanedimethanol divinyl ether; mixtures of C₁₄ and C₁₅ diol divinyl ethers; and mixtures of the above.

Preferred alkyl acrylate monomers include stearyl acrylate, lauryl acrylate and isodecyl acrylate. A particularly preferred one is lauryl acrylate. Preferred other hydrocarbon monomers include the C₁₀ dithiol, 1,10-decane dithiol, and the divinyl ether, hexanediol divinyl ether.

The monomer should be used in a quantity sufficient to adjust the total primary coating composition to a viscosity in the range of about 2,000 cps (centipoises) to 10,000 cps, measured by a Brookfield viscometer, model LVT, spindle speed of 6 rpm, spindle number 34, at 25°C.

C. The Photoinitiator

A third, optional, ingredient of the primary coating composition is a photoinitiator (C). In free radical systems, the photoinitiator is only required when ultraviolet radiation-cure is envisioned; in other embodiments, such as, for example, if electron beam cure of a free radical system is to be effected, the photoinitiator may be omitted. In cationically cured systems, however, a photoinitiator is required even when electron beam cure is to be effected.

The photoinitiator, when used in a small but effective amount to promote radiation cure, must provide reasonable cure speed without causing premature gelation of the composition. Further, it must not interfere with the optical clarity of the cured optical coating. Still further, the photoinitiator must itself be thermally stable, non-yellowing and efficient. The type of photoinitiator which is used will be dependent on whether a free radical-type system or a cationic cure type-system is used.

Suitable free radical-type photoinitiators include but are not limited to the following: hydroxycyclohexylphenyl ketone; hydroxymethylphenylpropanone; dimethoxyphenylacetophenone; 2-methyl-1-[4-(methyl thio)phenyl] -2-morpholinopropanone-1; 1-(4-isopropylphenyl)-2-hydroxy-2-methylpropan-1-one; 1-(4-dodecylphenyl)-2-hydroxy-2-methylpropan-1-one; 4-(2-hydroxyethoxy)phenyl-2(2-hydroxy-2-propyl)ketone; diethoxyphenyl acetophenone; and mixtures of these.

Suitable cationic cure-type photoinitiators include but are not limited to onium salts such as iodonium, sulfonium, arsonium, azonium, bromonium or selenonium, which are preferably chemically modified to render them more hydrophobic, e.g., by incorporating saturated hydrocarbon moieties such as alkyl or alkoxy substituents of from about 4 to about 18 carbon atoms. Particularly preferred cationic cure initiators include but are not limited to (4-octyloxyphenyl) phenyl iodonium hexafluoro antimonate; (4-octyloxyphenyl) diphenyl sulfonium hexafluoro antimonate; (4-decyloxyphenyl) phenyl iodonium hexafluoro antimonate; and (4-octadecyloxyphenyl) phenyl iodonium hexafluoro antimonate.

The photoinitiator, when used, preferably comprises from about 0.05 percent to about 10.0 percent by weight of the composition, based upon the total composition of the (A) through (C) ingredients. Preferably, the amount of photoinitiator is from about 0.2 percent to about 8.0 percent, and more preferably about 0.3 percent to about 7.0 percent by weight, based upon total weight of (A) (B) and (C) ingredients. Generally, cationic cured embodiments require lower amounts of photoinitiator than do free radical embodiments.

A particularly preferred free radical-type photoinitiator is hydroxycyclohexylphenyl ketone, and a particularly preferred cationic cure-type initiator is (4-octyloxyphenyl) phenyl iodonium hexafluoro antimonate. The photoinitiator should be chosen such that a cure speed, as measured in a dose versus modulus curve, of less than 1.0 J/cm², and preferably less than 0.5 J/cm², is required, when the photoinitiator is used in the designated amount.

D. The Optional Crosslinking Agent

Another optional ingredient is a crosslinking agent (D), which may be used in thiolene-type systems. This agent is typically a polyene or a polythiol which is capable of helping to coreact a multifunctional system. An example of a suitable crosslinking agent is trimethylethane trivinyl ether. When used, the crosslinking agent comprises from about 1 percent to about 50 percent by weight, and preferably from about 5 percent to about 30 percent by weight of the composition, based on the total weight of the (A), (B) and (C) components.

E. The Optional Stabilizer

To improve shelf life (storage stability) of the uncured coating, as well as to increase thermal and oxidative stability of the cured coating, one or more stabilizers may optionally be included in the composition. Examples of suitable stabilizers include but are not limited to tertiary amines such as diethylethanolamine, diethyl hydroxyl amine, and trihexylamine; hindered amines; organic phosphites; hindered phenols; mixtures thereof; and the like. Some particular examples of antioxidants which can be used include but are not limited to octadecyl-3-(3',5'-di-tert-butyl-4'-hydroxyphenyl) propionate; thiodiethylene bis (3,5-di-tert-butyl-4-hydroxy) hydrocinnamate; butylated paracresol-dicyclopentadiene copolymer; and tetrakis [methylene (3,5-di-tert-butyl-4-hydroxyhydrocinnamate)] methane. Additionally, certain silanes may be used in very small quantities, e.g., as low as about 0.0001 percent to about 0.01 percent by weight, as stabilizers. One example of a suitable silane stabilizer is 3-aminopropyltrimethoxy silane.

When a stabilizer is used, it may be incorporated in an amount from about 0.0001 percent to about 3.0 percent by weight, based on the weight of the (A) through (C) ingredients. Preferably, it is included in the range from about 0.25 percent to about 2.0 percent by weight, and more preferably in the range from about 0.5 percent to about 1.5 percent by weight, based on the total weight of the (A) through (C) ingredients. Desirable properties of a stabilizer include (1) non-migration (probably enhanced by low polarity and high molecular weight) and (2) basicity (to allow it to help in neutralizing residual acid which might prematurely initiate polymerization). Preferred stabilizers are octadecyl-3-(3',5'-di-tert-butyl-4'-hydroxyphenyl) propionate and trihexylamine.

20 F. The Optional Adhesion Promoter

Another optional ingredient is an adhesion promoter (F). Adhesion becomes a particularly pertinent problem in high humidity and high temperature environments, where delamination is more of a risk. Accordingly, in low humidity, low temperature situations (i.e., below about 50% relative humidity and about 25°C), the adhesion promoter is not required, although it may be required under conditions of higher humidity and higher temperature.

It is known in the art to use either acid-functional materials or organofunctional silanes to promote adhesion of resins to glass. While acid-functional materials are operative in some embodiments, organo-functional silanes are preferred. Acid-functional materials are less preferred because of their possible corrosivity towards the materials; their reactivity with certain systems (e.g., amin-enes); and their tendency to lose their adhesion properties on exposure to moisture. Silanes tend to be much more suitable in terms of these factors and, therefore, are the adhesion promoters of choice. Additionally, it is useful to have an adhesion promoter having a functionality which binds in with the system during cure, to maximize its adhesion promotion as well as to minimize the quantities of unbound volatiles. It is necessary to select a silane which does not react adversely with the chemistry of the system (e.g., amino-functional silanes in cationically cured systems).

Various suitable organofunctional silanes include but are not limited to amino-functional silanes; acrylamido-functional silanes; mercapto-functional silanes; allyl-functional silanes; vinyl-functional silanes; methacrylate-functional silanes; and acrylate-functional silanes. The adhesion promoters preferably are methoxy- or ethoxy-substituted as well.

Preferred organofunctional silanes include but are not limited to mercaptoalkyl trialkoxy silane, methacryloxyalkyl trialkoxy silane, aminoalkyl trialkoxy silane, vinyl trialkoxy silane, mixtures thereof, and the like.

The silane is, of course, chosen to be reactive with the rest of the system. Methacrylated silanes are desirable, when they are chosen to bind in well with the cured system, but tend to slow the cure speed of the system. The mercapto-functional adhesion promoters also chemically bind in during cure in many systems, but do not appreciably slow down the cure speed of the system. However, when mercapto-functional adhesion promoters are used in thiolene systems, it is necessary to adjust the stoichiometry of the polyene component accordingly. Allyl-functional and vinyl-functional silanes are also desirable when they are chosen to bind in well with the system used.

Some preferred organofunctional silanes that enhance adhesion in humid conditions include 3-aminopropyl triethoxy silane, 3-methacryloxy-propyltrimethoxy silane, gamma-mercaptopropyl trimethoxy silane, gamma-mercaptopropyl (gamma-mercaptopropyl) triethoxy silane, beta-(3,4-epoxycyclohexyl) ethyl trimethoxy silane, gamma-glycidoxypropyl trimethoxy silane, 3-vinylthiopropyl trimethoxy silane, vinyl-tris(beta-methoxyethoxy) silane, vinyl triacetoxy silane, and mixtures thereof. A particularly preferred adhesion promoter is gamma-mercaptopropyl trimethoxy silane.

The silane component, when used, should be incorporated into the composition in a small but effective amount to enhance the adhesion of the composition to the surface of an inorganic substrate, which in this embodiment, is glass, and in other embodiments, is glass, enamel, marble, granite or the like. The silane component, when used, comprises from about 0.1 percent to about 3.0 percent by weight of the composition, based on total weight of the ingredients (A), (B) and (C). Preferably, the silane comprises from about 0.2 percent to about 2.0 percent, and more preferably from about 0.3 percent to about 1.0 percent, based on the total weight of the oligomer, hydrocarbon monomer, and photoinitiator.

G. The Monomer Or Oligomer Containing An Aromatic Moiety

A further ingredient is a monomer or oligomer (G) which is capable of adjusting the refractive index of the composition. The refractive index of the formulation, in any case, should be 1.48 or higher. The monomer or oligomer contains at least (1) an aromatic moiety, (2) a moiety providing a reactive terminus, and (3) a hydrocarbon moiety. This compound is used in an amount effective for its stated purpose not in excess of about 60 percent by weight, and comprises from 5 percent to 60 percent by weight, preferably, from about 10 percent to about 40 percent by weight and more preferably from about 15 percent to about 30 percent by weight of the composition, based upon total weight of the ingredients (A), (B) and (C).

The monomer or oligomer is used to increase the refractive index for the reasons specified above. The aromatic moiety of the monomer or oligomer (B) is itself capable of raising the refractive index; however, the hydrocarbon moiety is required to increase the compatibility of this monomer with the hydrocarbon-based oligomer (A). The moiety containing the reactive terminus renders the compound compatible with the system as a whole, inasmuch as it has an available reactive group which allows it to crosslink with the rest of the composition upon ultraviolet curing, thus minimizing the volatiles content of the cured system.

Samples of aromatic monomers or oligomers additionally containing hydrocarbon character and a reactive group include but are not limited to polyalkylene glycol nonylphenylether acrylates such as polyethylene glycol nonylphenylether acrylate or polypropylene glycol nonylphenylether acrylate; polyalkylene glycol nonylphenylether methacrylates such as polyethylene glycol nonylphenylether methacrylate or polypropylene glycol nonylphenylether methacrylate; and mixtures of these. Refractive index modifiers containing termini other than acrylate or methacrylate are also suitable, so long as the terminus of this ingredient is coreactive with the rest of the system.

In each case, the phenyl group serves to increase the refractive index of the coating and the nonyl component renders the composition somewhat more hydrophobic and, therefore, compatible with the hydrophobic oligomer (A). The refractive index of the composition including this monomer or oligomer is higher than that of the composition comprising only (A), (B) and (C). A suitable primary coating composition should, again, have a refractive index of greater than or equal to about 1.48.

The refractive index of the primary coating must be higher than that of the cladding of the fiber. If the fibers coated with the coating composition of the present invention are down-doped, i.e., contain dopants which lower the refractive index of the fiber itself, the refractive index of the coating will be different enough from that of the fiber so that errant signals will be refracted away even without the incorporation of this component. Therefore, in such embodiments, the monomer or oligomer (G) is not essential to the composition.

H. Optional Chain Transfer Agents

Various other optional components may be used in the primary coating beyond the (A) through (G) components which are required or optional and which are described above. For example, optional chain transfer agents (H) may be used to control the modulus and glass transition temperature of the coating in acrylate or methacrylate systems. One way known in the art to control the molecular weight and, consequently, the modulus and glass transition temperature of a polymerization product is to use one or more chain transfer agents. It is postulated that the addition of a chain transfer agent to a formulation lowers the molecular weight of a polymer produced and results in a lower modulus, lower glass transition temperature coating.

Preferred chain transfer agents which may be used in acrylate or methacrylate systems are mercapto compounds, optionally having a hydrocarbon chain of at least eight carbon atoms. Examples of suitable mercapto chain transfer agents include methyl thioglycolate; methyl-3-mercaptopropionate; ethyl thioglycolate; butyl thioglycolate; butyl-3-mercaptopropionate; isoctyl thioglycolate; isoctyl-3-mercaptopropionate; isodecyl thioglycolate; isodecyl-3-mercaptopropionate; dodecyl thioglycolate; dodecyl-3-mercaptopropionate; octadecyl thioglycolate; and octadecyl-3-mercaptopropionate. Parathiocresol; thioglycolic acid; and 3-mercaptopropionic acid may also be used, but may display some incompatibility with the resin and may produce odor problems.

A particularly preferred chain transfer agent is octadecyl-3-mercaptopropionate (ODMP).

The chain transfer agent may, if used, comprise from about 0.1 percent to about 10.0 percent by weight of the composition based upon the total weight of ingredients (A) through (C). Preferably, the chain transfer agent comprises from about 0.25 percent to about 5.0 percent by weight, and still more preferably from about 0.5 percent to about 4.0 percent by weight based on the total weight of the (A), (B) and (C) components. A suitable chain transfer agent, when incorporated at about a 1.0 percent by weight level, lowers the glass transition temperature of the composition by about 3°C, lowers the modulus by about 689 kPa (100 psi), and apparently increases the adhesion of the coating to the glass fiber. When greater than about 2.0 percent by weight of the chain transfer agent is incorporated in the composition, the modulus and glass transition temperature of the composition are desirably lowered, but the shelf life is reduced somewhat.

One preferred primary composition for coating an optical fiber, then, comprises the following:

(A) from about 45 percent to about 65 percent by weight of an aliphatic urethane acrylate oligomer having as a backbone an hydrogenated 1,2-polybutadiene polymer;

(B) from about 10 percent to about 20 percent by weight of lauryl acrylate;

(C) from about 15 percent to about 30 percent by weight of polypropylene glycol nonylphenyl ether acrylate;

(D) from about 0.3 percent to about 1.0 percent by weight of gamma mercaptopropyl trimethoxy silane adhesion promoter;

(E) from about 2.0 percent to about 7.0 percent by weight of hydroxycyclohexylphenyl ketone photoinitiator;

(F) from about 0.5 percent to about 4.0 percent by weight of octadecyl-3-mercaptopropionate, a chain transfer agent; and

(G) from about 0.5 percent to about 1.5 percent by weight of octadecyl 3-(3',5'-di-tert-butyl-4'-hydroxyphenyl) propionate, wherein all of the stated percentages are percentages by weight, based upon total weight of (A) through (E), inclusive.

20 PREPARATION OF A COATED OPTICAL FIBER

The invention also relates to a process for preparing a coated optical fiber. The process comprises (i) applying to an optical glass fiber a primary coating layer comprising

(A) from about 10 percent to about 90 percent by weight, based upon total weight of (A), (B) and (C), of a reactively terminated urethane oligomer which is the reaction product of (i) a hydrocarbon polyol, the hydrocarbon portion of which has a molecular weight of from about 500 to about 4,000 (and preferably about 600 to 4,000); (ii) an aliphatic polyisocyanate; and (iii) an endcapping monomer capable of supplying a reactive terminus;

(B) from about 10 percent to about 90 percent by weight, based upon total weight of (A), (B) and (C), of a hydrocarbon monomer having 6 to 20 carbon atoms in its alkyl moiety and which is terminated with at least one end group capable of reacting with the reactive terminus of (A);

(C) optionally, from about 0.05 percent to about 10.0 percent by weight, based upon total weight of (A), (B) and (C), of a photoinitiator; and

(G) from 5% to 60% by weight of a monomer or oligomer selected from the group consisting of polyalkylene glycol nonylphenyl ether acrylates, polyalkylene glycol nonylphenyl ether methacrylates, and mixtures thereof;

(ii) applying atop said primary coating layer a secondary coating layer;

(iii) radiation-curing *in situ* said primary and secondary coating layers, e.g., by ultraviolet irradiation.

The primary and/or secondary coatings may be applied and cured by any method known in the art. A preferred method, whereby two coatings are applied wet-on-wet, is disclosed in U.S. Patent 4,474,830 to C. Taylor or AT&T Bell Laboratories. The coating or coatings may then be cured *in situ*, preferably by ultraviolet irradiation, to obtain a cured polymeric coating. Alternatively, the primary coating may be applied and cured, after which the secondary coating may be applied and cured.

COATED OPTICAL FIBERS OF THE INVENTION

The invention further relates to optical fibers, and especially to glass optical fibers, that are coated with the primary coating of the invention alone, or coated with both the primary coating and a secondary coating. These coatings may be prepared from reactively terminated materials that are radiation-curable and are radiation-cured *in situ* subsequent to application on glass fiber.

The primary coating surrounding the fiber forms a cured polymeric material preferably having a glass transition temperature (Tg) of no more than about -20°C and more preferably lower, i.e., about -30°C, or even lower.

The cured secondary coating should have a relatively high Tg, generally approximately 50°C, and a high tensile modulus, i.e., above 276 MPa (40,000 psi) and preferably about 689 MPa (100,000 psi). It is desirable for the secondary

coating to have a Tg higher than its highest use temperature. This is because at or near the Tg of the polymer, the physical properties change dramatically with small changes in temperature.

When cured, the coatings of the invention confer extraordinary thermal, hydrolytic and oxidative stability to the fiber and relieve stress thereon. Optical fibers having applied thereto the primary coatings of the invention and a secondary coating are highly moisture resistant and otherwise protected from environmental damage.

The coating compositions of the present invention have been disclosed hereinabove for use as optical fiber coatings. However, it is to be understood that these coatings may be used in any embodiment wherein stable, moisture resistant coatings are desired, especially for coating the surface of an optically useful article. For example, the coatings may be used for such diverse end uses as coating sheet glass (i.e., in the preparation of safety glass) to coating vinyl materials (e.g., in preparing no-wax floors). Other optically useful articles which may be prepared include, but are not limited to, photoswitches, photorelay devices, microelectronic devices, photocoupling devices, and so forth.

EXAMPLES

The following Examples serve to further illustrate the invention. In these Examples and elsewhere throughout this application, all parts and percentages are by weight, on a dry solids basis, and all temperatures are in degrees centigrade unless expressly stated to be otherwise. In all of the examples, cure speeds were measured with an International Light IL 745-A radiometer with model A309 light bug. Unlike elsewhere in the application, where percentage by weight referred to the total weight of either the (A) through (C) ingredients for the primary coating, parts by weight in the Examples refers to the total composition described in that Example, including all components. The optional ingredients are identified by an asterisk (*) in the Examples. The other components are essential for use, if the exemplified coating is to meet the rigorous requirements for a commercially acceptable coating for optical glass fiber. It should be noted that, although the photoinitiator is identified as optional in the specification, it is, in fact, required in any system cured by ultraviolet irradiation, and therefore is not denoted as optional in such Examples. It should also be noted that, unless otherwise indicated, throughout the Examples and the remainder of this application, "modulus" refers to 2.5% tensile modulus, measured using an Instron tensile tester.

Example I

Primary Coating for Optical Fibers

An ultraviolet-curable, primary coating composition was made up as follows:

Ingredient	Parts by Weight
aliphatic urethane acrylate oligomer with saturated hydrocarbon backbone (A)	55.00
lauryl acrylate (B)	14.50
hydroxycyclohexylphenyl ketone photoinitiator (C)	6.00
octyldecyl-3-(3',5'-di-tert-butyl-4'-hydroxy) hydrocinnamate stabilizer (E)*	1.50
octadecyl-3-mercaptopropionate chain transfer agent (H)*	2.75
gamma-mercaptopropyl trimethoxy silane adhesion promoter (F)*	0.75
polypropylene glycol nonylphenylether acrylate (G)	19.50

The viscosity of the uncured coating was 4.005 Pas (4005 cps) (at 25°C using a Brookfield viscometer, model LVT, 6 rpm, #34 spindle).

A 0.152mm (six mil) coating of this composition was applied to a flat glass sheet using a Bird applicator and cured in air at 0.7 J/cm² using a 7.87 watts per mm (200 watts per inch) medium pressure mercury vapor lamp.

The cured primary coating which resulted had a glass transition temperature of about -39.6°C, and a cured film refractive index of 1.492. The effect of temperature on 2.5% tensile modulus was as follows: 25°C: 1.05 MPa (153.0 psi); 0°C: 1.16 MPa (167.8 psi); -20°C: 5.96 MPa (864.7 psi); -40°C: 343 MPa (49,807 psi); -60°C 1371 MPa (199,018 psi). The cure speed, as determined by a modulus versus dose curve, was determined to be about 0.3 J/cm².

Water absorption of the sample was measured as follows. The cured film was equilibrated at 50% ($\pm 5\%$) relative humidity and 23°C ($\pm 2\%$) for 48 hours. The sample was weighed and a weight "A" recorded. The sample was then soaked for 24 hours at 25°C in distilled water, then patted dry and weighed. This weight was recorded as "B". The sam-

ple was next placed in a vacuum oven under 1.33 kPa (10mm Hg) pressure at 25°C for 24 hours, removed, and again equilibrated at 50% ($\pm 5\%$) relative humidity at 23°C ($\pm 2^\circ\text{C}$) for 48 hours and weighed. This third weight was recorded as "C". Percent water absorption measured as

5

$$\frac{B-C}{A} \times 100\%$$

was about 1.48%.

10 Other samples of this 6 mil thick coating were tested as follows to determine the effect of accelerated aging on various properties. The samples were equilibrated at 50% ($\pm 5\%$) relative humidity and 23°C ($\pm 2^\circ\text{C}$) for 48 hours and then placed on clips and hung in an environmental chamber under the following accelerated aging conditions:

- 15 (1) 125°C for 7 days;
 (2) 93.3°C for 10 days;
 (3) 93.3°C for 30 days;
 (4) 93.3°C, 95% relative humidity, for 10 days; and
 (5) 93.3°C, 95% relative humidity, for 30 days.
- 20 After the designated time period, the samples were removed and again equilibrated for 48 hours. The following properties were recorded. These data indicate excellent stability against thermal and hydrolytic stress.

	125°C, 7 days	93.3°C, 10 days	93.3°C, 30 days	93.3°C, 95% relative humidity, 10 days	93.3°C, 95% relative humidity, 30 days
weight change	-7.68%	-6.71%	-6.98%	-0.71%	-1.62%
Tg midpoint	-34.6°C	-34.6°C	-36.6°C	-35.5°C	-36.6°C
Instron (tensile) modulus change (at 25°C)	+33.1%	+39.4%	+37.9%	+27.9%	+19.5%

35 **Example II**

Another Primary Coating for Optical Fiber

An ultraviolet radiation-curable primary coating was made up as follows:

40

Ingredient	Parts by Weight
aliphatic urethane acrylate oligomer with saturated hydrocarbon backbone (A)	56.00
isodecyl acrylate (B)	11.75
hydroxymethylphenyl-propanone photoinitiator (C)	6.00
thiodiethylene bis (3,5-di-tert-butyl-4-hydroxy) hydrocinnamate stabilizer (E)*	0.50
gamma mercaptopropyl trimethoxy silane adhesion promoter (F)*	0.75
polyethylene glycol nonylphenyl ether acrylate (G)	25.00

The above formulation used several different ingredients, and somewhat different proportions, than were used in Example I. This formulation has a viscosity of 5.65 Pas (5650 cps) at 25°C using a Brookfield viscometer, model LVT, 6 rpm, #34 spindle, and a refractive index of 1.4849 at 21.8°C. When coated onto a glass plate in the same manner as in Example I and cured in air at 0.7 J/cm² under a medium pressure 7.87 watts per mm (200 watts per inch) mercury vapor lamp, the resultant 0.152 mm (6 mil) thickness coating had a tensile modulus of 2.04 MPa (295.8 psi), a Tg of -28.7°C,

good adhesion to glass and low water absorption (2.02%). The coating performed well in high humidity and dry accelerated aging tests.

Example III

A Further Primary Coating for Optical Fiber

The following primary coating was formulated:

Ingredient	Parts by Weight
aliphatic urethane acrylate oligomer with saturated hydrocarbon backbone (A)	56.00
octyl/decyl acrylate (B)	11.75
hydroxymethylphenyl-propanone photoinitiator (C)	6.00
thiodiethylene bis (3', 5'-di-tert-butyl-4-hydroxy) hydrocinnamate stabilizer (E)*	0.50
gamma mercaptopropyl trimethoxy silane adhesion promoter (F)*	0.75
polyethylene glycol nonylphenylether acrylate (lower MW polyethylene glycol than that of Example II) (G)	25.00

This formulation demonstrated the use of alternate (B) and (C) materials. This formulation had a viscosity of 3.89

Pas (3890 cps) at 25°C using a Brookfield viscometer, model LVT, spindle #34, 6 rpm. When applied to a glass plate as in Example I and cured in air at 0.7 J/cm² under 7.87 watts per mm (200 watts per inch) medium pressure mercury vapor lamp, it had a tensile modulus of 1.90 MPa (276.0 psi), good adhesion to glass, and low water absorption (1.89%).

Example IV

Another Primary Coating for Optical Fibers

The following primary coating was formulated:

Ingredient	Parts by Weight
aliphatic urethane acrylate oligomer with saturated hydrocarbon backbone (A)	64.00
isodecyl acrylate (B)	20.50
hydroxymethylphenyl-propanone photoinitiator (C)	6.00
thiodiethylene bis (3', 5'-di-tert-butyl-4'-hydroxy) hydrocinnamate stabilizer (E)*	0.50
gamma mercaptopropyl trimethoxy silane adhesion promoter (F)*	0.35
polyethylene glycol nonylphenylether acrylate (same MW as in Example III) (G)	8.00
isoctyl-3-mercaptopropionate chain transfer agent (H)*	0.65

This formulation demonstrated the use of different proportions of ingredients, the use of an alternate chain transfer agent, and the use of a low concentration of the component (G) to adjust the refractive index. The formulation had a

viscosity of 5.95 Pas (5950 cps) at 25°C using a Brookfield viscometer, model LVT, 6 rpm, #34 spindle, and a refractive index of 1.4796 at 21.5°C. When applied to a glass sheet as a 0.152 mm (6 mil) coating as in Example I and cured in air at 0.7 J/cm² under a medium pressure 7.87 watts per mm (200 watts per inch) mercury vapor lamp, it had a tensile modulus of 1.93 MPa (280 psi), a Tg of -31.2°C, good adhesion to glass, and low water absorption (1.41 %).

Example V**A Slower Curing Primary Coating for Optical Fibers**

5 A radiation curable primary coating was formulated as follows:

Ingredient	Parts by Weight
aliphatic urethane acrylate oligomer with saturated hydrocarbon backbone (A)	54.00
lauryl acrylate (B)	14.00
hydroxycyclohexylphenyl ketone photoinitiator (C)	6.00
octadecyl-3-mercaptopropionate chain transfer agent (H)*	4.00
octadecyl-3(3',5'-di-tert-butyl-4'-hydroxy phenyl) propionate stabilizer (E)*	1.50
gamma-methacryloxypropyl trimethoxysilane adhesion promoter (F)*	1.00
polypropylene glycol nonylphenylether acrylate (G)	19.50

20 The viscosity of the uncured coating was 3.75 Pas (3750 cps) (at 25°C using a Brookfield viscometer, model LVT, 6 rpm, #34 spindle).

When coated on a glass sheet at 0.152 mm (6 mils) and cured as in previous Examples, the coating adhered well under high humidity conditions but cured much slower than the coating of Example I. Similar formulations can be prepared that have much more rapid rates of cure, by substituting acrylated silanes for the methacrylated silane used in this Example.

The cured primary coating which resulted had a glass transition temperature of about -39.4°C, a tensile modulus of 1.07 MPa (155.5 psi), and a cured film refractive index of 1.492. The cure speed was determined to be 0.5 J/cm², and the water absorption was 1.40%, both being determined according to the methods of Example I.

30 Weight change, glass transition temperature and Instron (tensile) modulus change were measured under the same conditions as in Example I and recorded to be as follows. Again, excellent hydrolytic and thermal stability was observed.

	125°C, 7 days	93.3°C, 10 days	93.3°C, 30 days	93.3°C, 95% relative humidity, 10 days	93.3°C, 95% relative humidity, 30 days
weight change	-7.94%	-6.84%	-7.19%	-1.97%	-1.58%
Tg midpoint	-35.9°C	-36.1°C	-35.5°C	-35.7°C	-36.4°C
Instron tensile modulus change (at 25°C)	+33.8%	+36.1%	+36.0%	+20.1%	+10.9%

CONCLUSION

Optical glass fibers coated with the primary and secondary coatings of the present invention have been designed to possess several important qualities rendering the coated fibers useful for many applications and particularly suitable in local area networks for fiber-to-the-home uses.

50 It has now been discovered that virtually any chemistry which is capable of reacting together certain specific functional backbones and which does not adversely effect the properties of the composition when cured, may be used to formulate a composition possessing these qualities.

55 The primary coatings, when cured, possess a refractive index greater than that of the underlying fiber (i.e., about 1.48), a tensile modulus of less than about 3.45 MPa (500 psi) and a glass transition temperature of less than about -20°C, and preferably less than about -30°C. The primary coatings are sufficiently hydrophobic to resist moisture penetration and consequent hydrolysis of the underlying optical fibers; have a sufficiently low tensile modulus to cushion and protect the fibers; and a sufficiently low glass transition temperature in the use range of the fibers.

The primary coatings have suitable viscosities for coating and, when cured, contain minimal quantities of unbound

reactants (volatiles) and have good shelf life characteristics.

The present invention permits the production of coating formulations meeting these criteria, and the manufacture of coated optical fibers well suited for use in fiber-to-the-home applications, though there are many uses for coatings that are prepared in accordance with the invention, that do not have to meet the stringent requirements imposed on optical fiber coatings.

It has now been discovered that an optimized composition can be formulated having the above properties by using the backbones of the components described herein and in the parent application, but that any end groups capable of reacting, via free radical initiation, cationic cure or other method, may be employed. It was also discovered that, in connection with the present invention, a coating formulated in the absence of a silane adhesion promoter and a refractive index modifier, and, in some cases, a photoinitiator, all believed in the parent application to be essential in all embodiments, could in some instances be satisfactory.

Thus, useful coatings for many purposes may be formulated, the primary coating comprising:

(A) from 10% to 90% by weight of a reactively terminated urethane oligomer which is the reaction product of (i) a hydrocarbon polyol, the hydrocarbon portion of which is from 500 to 4,000 molecular weight; (ii) an aliphatic polyisocyanate; and (iii) an endcapping monomer capable of supplying a reactive terminus;

(B) from 10% to 90% by weight of a hydrocarbon monomer containing from 6 to 20 carbon atoms which is terminated with at least one end group capable of reacting with the reactive terminus of (A);

(C) optionally, from 0.5% to 10.0% by weight of photoinitiator; and

(G) from 5% to 60% by weight of a monomer or oligomer selected from the group consisting of polyalkylene glycol nonylphenyl ether acrylates, polyalkylene glycol nonylphenyl ether methacrylates, and mixtures thereof, all of the stated percentages are percentages by weight based on the total weight of (A), (B) and (C), wherein the coating, when cured, has a refractive index of 1.48 or higher

with one or more of the following optional components:

(D) a crosslinking agent capable of helping to coreact multifunctional systems;

(E) a stabilizer or antioxidant that imparts added shelf life and storage stability to the coating composition;

(F) an organofunctional silane adhesion promoter capable of preventing delamination under conditions of high humidity and high temperature; and

(H) a mercapto functional chain transfer agent.

The combination of (B) and (C) with (A) has the effect of producing a coating of mutually compatible ingredients, that is radiation-curable.

In preferred embodiments, the (A) component may be acrylate, thiol or vinyl ether terminated.

The composition may be, in one embodiment, used as a primary coating for an optical fiber, the coating having a cured tensile modulus of less than 3.45 MPa (500 psi) (at room temperature, based on a 0.152mm (6 mil) thick film).

In other embodiments, compositions of this kind can be formulated to be useful as interlayers for laminated safety glass, and as coatings for electronic devices such as photocells and photoswitches, for example.

Claims

1. A radiation-curable coating for the surface of an optically useful article comprising

(A) from 10% to 90% by weight of a reactively terminated urethane oligomer which is the reaction product of (i) a hydrocarbon polyol, the hydrocarbon portion of which is from 500 to 4,000 molecular weight; (ii) an aliphatic polyisocyanate; and (iii) an endcapping monomer capable of supplying a reactive terminus;

(B) from 10% to 90% by weight of a hydrocarbon monomer containing from 6 to 20 carbon atoms which is terminated with at least one end group capable of reacting with the reactive terminus of (A);

(C) optionally, from 0.05% to 10.0% by weight of photoinitiator; and

(G) from 5% to 60% by weight of a monomer or oligomer selected from the group consisting of polyalkylene glycol nonylphenyl ether acrylates, polyalkylene glycol nonylphenyl ether methacrylates, and mixtures thereof, all of the stated percentages are percentages by weight based on the total weight of (A), (B) and (C), wherein the coating, when cured, has a refractive index of 1.48 or higher.

2. A radiation-curable coating according to claim 1, wherein said oligomer (A) is characterised by the formula



5

where

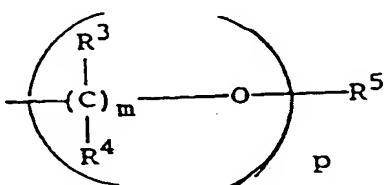
10 R¹ is a linear or branched hydrocarbon polymer of from 500 to 4,000 molecular weight selected from the group consisting of fully or partially hydrogenated 1,2-polybutadiene; 1,2-polybutadiene hydrogenated to an iodine number of from 9 to 21; fully or partially hydrogenated polyisobutylene; polybutene polyol; and hydrogenated dimer diol;

R² is a linear, branched or cyclic alkylene of from six to twenty carbon atoms; and

15 X and Q are independently either

(a) a radical of the formula:

20



25

wherein R⁵ is selected from the group consisting of acrylic, methacrylic, vinylic, allylic, itaconic, crotonic, styrenic, acrylamide, norbornenyl, acetylenic, epoxy, mercapto and amino; R³ and R⁴ are independently hydrogen, methyl, ethyl or propyl; m is an integer from 1 to 10; and p is either zero or one,

(b) a saturated alkyl radical of from nine to twenty carbon atoms, with the proviso that said oligomer must possess at least one reactive terminal group.

- 35 3. A radiation-curable coating according to claim 1 or claim 2, wherein said monomer (B) is selected from the group consisting of hexyl acrylate; hexyl methacrylate; 2-ethylhexyl acrylate; 2-ethylhexyl methacrylate; octyl acrylate; octyl methacrylate; isoctyl acrylate; isoctyl methacrylate; decyl acrylate; decyl methacrylate; isodecyl acrylate; isodecyl methacrylate; lauryl acrylate; lauryl methacrylate; stearyl acrylate; stearyl methacrylate; C₁₄-C₁₅ hydrocarbon diol diacrylates; C₁₄-C₁₅ hydrocarbon diol dimethacrylates; tridecyl acrylate; tridecyl methacrylate; palmitic acrylate; palmitic methacrylate; 1,10-decane dithiol; hexanediol divinyl ether; mercaptoacetate esters; mercapto-propionate esters; lauryl vinyl ether; ethylhexylvinyl ether; isodecyl vinyl ether; butanediol divinyl ether; cyclohexanedimethanol divinyl ether; C₁₄-C₁₅ diol divinyl ethers; and mixtures thereof.

40 4. A radiation-curable coating according to any one of claims 1 to 3, additionally comprising from 1.0% to 50.0% by weight, based on total weight of (A), (B) and (C), of a crosslinking agent (D).

45 5. A radiation-curable coating according to claim 4, wherein said crosslinking agent (D) is trimethylol ethane trivinyl ether.

50 6. A radiation-curable coating according to any one of claims 1 to 5, additionally comprising from 0.0001% to 3.0% by weight, based on total weight of (A), (B) and (C), of a stabiliser (E).

55 7. A radiation-curable coating according to claim 6, wherein said stabiliser (E) is selected from the group consisting of diethyl hydroxylamine; organic phosphites; hindered phenols; trihexylamine; octadecyl-3-(3',5'-di-tert-butyl-4'-hydroxy phenyl) propionate; thiobis(2-tert-butyl-4-hydroxy) hydrocinnamate; butylated paracresol-dicyclopentadiene copolymer; tetrakis [methylene (3,5-di-tert-butyl-4-hydroxyhydrocinnamate)] methane; 3-amino-propyltrimethoxy silane; and mixtures thereof.

8. A radiation-curable coating according to any one of claims 1 to 7, additionally comprising from 0.1% to 3.0% by weight, based on total weight of (A), (B) and (C), of an organofunctional silane adhesion promoter (F).
- 5 9. A radiation-curable coating according to any one of claims 1 to 8, wherein said photoinitiator (C) is selected from the group consisting of hydroxymethylphenyl propanone; hydroxycyclohexylphenyl ketone; dimethoxyphenyl acetophenone; 2-methyl-1,4-(methyl thio) phenyl-2-morpholino-propane-1; 1-(4-isopropyl-phenyl)-2-hydroxy-2-methylpropan-1-one; 1-(4-dodecylphenyl)-2-hydroxy-2-methylpropan-1-one; 4-(2-hydroxyethoxy)phenyl-2(2-hydroxy-2-propyl)ketone; diethoxyacetophenone; 2,2-di-sec-butoxyacetophenone; diethoxyphenyl acetophenone; (4-octyloxyphenyl) phenyl iodonium hexafluoro antimonate; (4-octyloxyphenyl) diphenyl sulfonium hexafluoro antimonate; (4-decyloxyphenyl) phenyl iodonium hexafluoro antimonate; (4-octadecyloxyphenyl) phenyl iodonium hexafluoro antimonate; and mixtures thereof.
- 10 10. A radiation-curable coating according to any one of claims 1 to 9, additionally comprising from 0.1 % to 10% by weight, based on total weight of (A), (B) and (C), of a mercapto functional chain transfer agent (H).
- 15 11. A radiation-curable coating according to any one of claims 1 to 10, which is a primary coating for an optical fiber.
12. A radiation-curable coating according to claim 1, which is a primary coating for an optical fiber, wherein the following components are present:
- 20 (A) from 30% to 70% by weight of an aliphatic vinyl ether urethane oligomer having as a backbone a hydrogenated 1,2-polybutadiene polymer;
- 25 (B) from 10% to 40% by weight of 1,10-decane dithiol;
- (C) from 2% to 10% by weight of hydroxymethyl phenyl propanone photoinitiator;
- (D) from 1% to 20% by weight of trimethylol ethane trivinyl ether crosslinking agent; and
- 30 (E) from 0.001% to 0.1% by weight of diethyl hydroxyl amine stabiliser,
- and wherein all of the stated percentages are percentages by weight based upon total weight of (A), (B) and (C).
- 35 13. A radiation-curable coating according to claim 1, which is a primary coating for an optical fiber, wherein the following components are present:
- 40 (A) from 20% to 80% by weight of an aliphatic vinyl ether-terminated urethane oligomer having as a backbone an hydrogenated 1,2-polybutadiene polymer;
- (B) from 10% to 80% by weight of hexanediol divinyl ether; and
- 45 (C) from 0.05% to 5.0% by weight of (4-octyloxyphenyl) phenyl iodonium hexafluoro antimonate cationic cure initiator,
- and wherein all of the stated percentages are percentages by weight based upon total weight of (A), (B) and (C).
- 50 14. A coated optical fiber comprising an optical fiber and a primary coating layer according to any one of claims 1 to 13.
15. A coated optical fiber according to claim 14, wherein said primary coating layer was radiation cured subsequent to application on said glass fiber.
- 55 16. A coated optical fiber according to claim 14 wherein said coating layer is formed from reactively terminated materials that are radiation-curable and that are radiation-cured *in situ*.
17. A process for preparing a coated optical fiber comprising
- (i) applying to an optical fiber a primary coating layer comprising a composition according to any one of claims 1 to 13; and

- (ii) applying atop said primary coating layer a secondary coating layer; and
- (iii) radiation-curing in situ said primary and secondary coating layers.

5 18. A process according to claim 17, wherein the primary coating layer comprises

- (A) from 45% to 65% by weight of an aliphatic urethane acrylate oligomer having as a backbone an hydrogenated 1,2-polybutadiene polymer;
- 10 (B) from 10% to 20% by weight of lauryl acrylate;
- (C) from 15% to 30% by weight of polypropylene glycol nonylphenyl ether acrylate;
- 15 (D) from 0.3% to 1.0% by weight of gamma mercaptopropyl trimethoxy silane adhesion promoter; and
- (E) from 2.0% to 7.0% by weight of hydroxycyclohexylphenyl ketone photoinitiator,

wherein all of the stated percentages are percentages by weight based upon total weight of (A), (B) and (C).

20 Patentansprüche

1. Strahlungshärtbarer Überzug für die Oberfläche eines für optische Zwecke geeigneten Gegenstandes, umfassend

- (A) 10 bis 90 Gew.-% eines reaktiv terminierten Urethanoligomers, das das Reaktionsprodukt ist von (i) einem Kohlenwasserstoffpolyol, wobei der Kohlenwasserstoffanteil davon ein Molekulargewicht von 500 bis 4 000 aufweist, (ii) einem aliphatischen Polyisocyanat und (iii) einem Endkappenmonomer, das in der Lage ist, einen reaktiven Terminus bereitzustellen,
- 25 (B) 10 bis 90 Gew.-% eines Kohlenwasserstoffmonomers, enthaltend 6 bis 20 Kohlenstoffatome, das mit mindestens einer Endgruppe terminiert ist, die mit dem reaktiven Terminus von (A) reagieren kann,
- 30 (C) gegebenenfalls 0,05 bis 10,0 Gew.-% eines Photoinitiatoren, und
- (G) 5 bis 60 Gew.-% eines Monomers oder Oligomers, ausgewählt aus der Gruppe, bestehend aus Polyalkylenglykol-nonylphenyletheracrylaten, Polyalkylenglykol-nonylphenylethermethacrylaten und Gemischen davon,
35 wobei alle angegebenen Prozentsätze Gewichtsprozent sind, bezogen auf das Gesamtgewicht von (A), (B) und (C),
wobei der Überzug, wenn er gehärtet ist, einen Brechungsindex von 1,48 oder mehr aufweist.

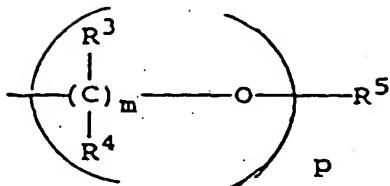
40 2. Strahlungshärtbarer Überzug nach Anspruch 1, wobei das Oligomer (A) charakterisiert ist durch die Formel



45 wobei

- R¹ ein lineares oder verzweigtes Kohlenwasserstoffpolymer mit einem Molekulargewicht von 500 bis 4 000 ist, ausgewählt aus der Gruppe, bestehend aus vollständig oder teilweise hydriertem 1,2-Polybutadien; 1,2-Polybutadien, das bis zu einer Iodzahl von 9 bis 21 hydriert ist; vollständig oder teilweise hydriertem Polyisobutylene; Polybutenpolyol und hydriertem dimeren Diol;
- R² ein lineares, verzweigtes oder cyclisches Alkylen mit sechs bis zwanzig Kohlenstoffatomen ist, und X und Q unabhängig

55 entweder (a) ein Rest der Formel:



sind, wobei R^5 ausgewählt ist aus der Gruppe, bestehend aus Acryl, Methacryl, Vinyl, Allyl, Itaconyl, Crotonyl, Styryl, Acrylamid, Norbornenyl, Acetylenyl, Epoxy, Mercapto und Amino, R^3 und R^4 unabhängig Wasserstoff, Methyl, Ethyl oder Propyl sind, m eine ganze Zahl von 1 bis 10 ist und p entweder null oder eins ist, oder

(b) ein gesättigter Alkylrest mit neun bis zwanzig Kohlenstoffatomen sind, mit der Maßgabe, daß das Oligomer mindestens eine reaktive terminale Gruppe enthalten muß.

- 20 3. Strahlungshärtbarer Überzug nach Anspruch 1 oder 2, wobei das Monomer (B) ausgewählt ist aus der Gruppe, bestehend aus Hexyl-acrylat, Hexyl-methacrylat, 2-Ethylhexyl-acrylat, 2-Ethylhexyl-methacrylat, Octyl-acrylat, Octyl-methacrylat, Isooctyl-acrylat, Isooctyl-methacrylat, Decyl-acrylat, Decyl-methacrylat, Isodecyl-acrylat, Isodecyl-methacrylat, Lauryl-acrylat, Lauryl-methacrylat, Stearyl-acrylat, Stearyl-methacrylat, $\text{C}_{14}\text{-C}_{15}$ -Kohlenwasserstoff-diol-diacrylaten, $\text{C}_{14}\text{-C}_{15}$ -Kohlenwasserstoff-diol-dimethacrylaten, Tridecyl-acrylat, Tridecyl-methacrylat, Palmityl-acrylat, Palmityl-methacrylat, 1,10-Decan-dithiol, Hexan-diol-divinylether, Mercaptacetat-estern, Mercaptopropionat-estern, Lauryl-vinyl-ether, Ethylhexylvinyl-ether, Isodecyl-vinyl-ether, Butandiol-divinyl-ether, Cyclohexandimethanol-divinyl-ether, $\text{C}_{14}\text{-C}_{15}$ -Diol-divinyl-ethers und Gemischen davon.
- 30 4. Strahlungshärtbarer Überzug nach einem der Ansprüche 1 bis 3, enthaltend zusätzlich 1,0 bis 50,0 Gew.-%, bezogen auf das Gesamtgewicht von (A), (B) und (C), eines Vernetzungsmittels (D).
5. Strahlungshärtbarer Überzug nach Anspruch 4, wobei das Vernetzungsmittel (D) Trimethylol-ethan-trivinyl-ether ist.
- 35 6. Strahlungshärtbarer Überzug nach einem der Ansprüche 1 bis 5, umfassend zusätzlich 0,0001 bis 3,0 Gew.-%, bezogen auf das Gesamtgewicht von (A), (B) und (C), eines Stabilisators (E).
7. Strahlungshärtbarer Überzug nach Anspruch 6, wobei der Stabilisator (E) ausgewählt ist aus der Gruppe, bestehend aus Diethylhydroxylamin, organischen Phosphiten, gehinderten Phenolen, Trihexylamin, Octadecyl-3-(3',5'-di-tert-butyl-4'-hydroxy-phenyl)-propionat, Thiodiethylen-bis-(3,5-di-tert-butyl-4-hydroxy)hydrocinnamat, butyliertem p-Kresoldicyclopentadien-Copolymer, Tetrakis-[methylen-(3,5-di-tert-butyl-4-hydroxyhydrocinnamat)]methan, 3-Aminopropyltrimethoxysilan und Gemischen davon.
- 40 8. Strahlungshärtbarer Überzug nach einem der Ansprüche 1 bis 7, enthaltend zusätzlich 0,1 bis 3,0 Gew.-%, bezogen auf das Gesamtgewicht von (A), (B) und (C), eines organofunktionellen Silan-Haftverstärkers (F).
9. Strahlungshärtbarer Überzug nach einem der Ansprüche 1 bis 8, wobei der Photoinitiator (C) ausgewählt ist aus der Gruppe, bestehend aus Hydroxymethylphenyl-propanon, Hydroxycyclohexylphenylketon, Dimethoxyphenylacetophenon, 2-Methyl-1,4-(methylthio)-phenyl-2-morpholinopropan-1; 1-(4-Isopropyl-phenyl)-2-hydroxy-2-methylpropan-1-on; 1-(4-Dodecylphenyl)-2-hydroxy-2-methylpropan-1-on; 4-(2-Hydroxyethoxy)phenyl-2(2-hydroxy-2-propyl)keton; Diethoxyacetophenon; 2,2-Di-sec-butoxyacetophenon; Diethoxyphenylacetophenon; (4-Octyloxy-phenyl)phenyl-iodonium-hexafluor-antimonat; (4-Octyloxyphenyl)-diphenyl-sulfonium-hexafluorantimonat; (4-Decyloxyphenyl)phenyl-iodonium-hexafluor-antimonat; (4-Octadecyloxyphenyl)-phenyl-iodonium-hexafluor-antimonat und Gemischen davon.
- 50 10. Strahlungshärtbarer Überzug nach einem der Ansprüche 1 bis 9, enthaltend zusätzlich 0,1 bis 10 Gew.-%, bezogen auf das Gesamtgewicht von (A), (B) und (C), eines mercaptofunktionellen Kettenübertragungsmittels (H).

11. Strahlungshärtbarer Überzug nach einem der Ansprüche 1 bis 10, der ein erster Überzug für eine optische Faser ist.

12. Strahlungshärtbarer Überzug nach Anspruch 1, der ein erster Überzug für eine optische Faser ist, wobei die folgenden Bestandteile vorhanden sind:

(A) 30 bis 70 Gew.-% eines aliphatischen Vinyletherurethan-Oligomers mit einem hydrierten 1,2-Polybutadienpolymer als Hauptkette,

(B) 10 bis 40 Gew.-% 1,10-Decan-dithiol,

(C) 2 bis 10 Gew.-% Hydroxymethylphenylpropanon-Photoinitiator,

(D) 1 bis 20 Gew.-% Trimethylolethantrivinylether-Vernetzungsmittel, und

(E) 0,001 bis 0,1 Gew.-% Diethylhydroxylamin-Stabilisator, und wobei alle angegebenen Prozentsätze Gewichtsprozent sind, bezogen auf das Gesamtgewicht von (A), (B) und (C).

13. Strahlungshärtbarer Überzug nach Anspruch 1, der ein erster Überzug für eine optische Faser ist, wobei die folgenden Bestandteile vorhanden sind:

(A) 20 bis 80 Gew.-% eines aliphatischen Vinylether-terminierten Urethanoligomers mit einem hydrierten 1,2-Polybutadienpolymer als Hauptkette,

(B) 10 bis 80 Gew.-% Hexandioldivinylether, und

(C) 0,05 bis 5,0 Gew.-% (4-Octyloxyphenyl)phenyliodoniumhexafluorantimonat als kationischer Härtungsinitiator,

und wobei alle angegebenen Prozentsätze Gewichtsprozent sind, bezogen auf das Gesamtgewicht von (A), (B) und (C).

14. Beschichtete optische Faser, umfassend eine optische Faser und eine erste Überzugsschicht nach einem der Ansprüche 1 bis 13.

15. Beschichtete optische Faser nach Anspruch 14, wobei die erste Überzugsschicht nach dem Aufbringen auf die Glasfaser durch Strahlung gehärtet worden ist.

16. Beschichtete optische Faser nach Anspruch 14, wobei die Überzugsschicht aus reaktiv terminierten Materialien, die strahlungshärtbar sind und in situ durch Strahlung gehärtet worden sind, gebildet ist.

17. Verfahren zur Herstellung einer beschichteten optischen Faser, umfassend

(i) Aufbringen einer ersten Überzugsschicht, umfassend eine Masse nach einem der Ansprüche 1 bis 13, auf eine optische Faser und

(ii) Aufbringen einer zweiten Überzugsschicht auf die erste Überzugsschicht und

(iii) Strahlungshärten der ersten Überzugsschicht und der zweiten Überzugsschicht in situ.

18. Verfahren nach Anspruch 17, wobei die erste Überzugsschicht umfaßt:

(A) 45 bis 65 Gew.-% eines aliphatischen Urethanacrylat-Oligomers mit einem hydrierten 1,2-Polybutadienpolymer als Hauptkette,

(B) 10 bis 20 Gew.-% Laurylacrylat,

(C) 15 bis 30 Gew.-% Polypropylenglykolnonylphenyletheracrylat,

(D) 0,3 bis 1,0 Gew.-% γ -Mercaptopropyltrimethoxsilan-Haftverstrker und

(E) 2,0 bis 7,0 Gew.-% Hydroxycyclohexylphenylketon-Photoinitiator,

5 wobei alle angegebenen Prozentstze Gewichtsprozent sind, bezogen auf das Gesamtgewicht von (A), (B) und (C).

Revendications

10 1. Revtement durcissable par rayonnement pour la surface d'un article optiquement utilisable, comprenant :

(A) de 10 %  90 % en poids d'un oligomre d'urthane ayant un groupe terminal ractif, qui est le produit de
raction de (i) un polyol hydrocarbon dont la partie hydrocarbone a un poids molculaire de 500  4 000, (ii)
15 un polyisocyanate aliphatique, et (iii) un monomre de recouvrement terminal capable de fournir un groupe terminal apte  ragir ;

(B) de 10 %  90 % en poids d'un monomre hydrocarbon contenant de 6  20 atomes de carbone, qui est termin par au moins un groupe terminal capable de ragir avec le groupe terminal ractif de (A);

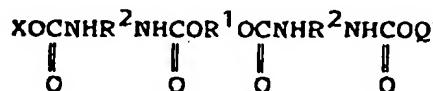
(C) ventuellement, de 0,05 %  10,0 % en poids de photo-initiateur ; et

20 (G) de 5 %  60 % en poids d'un monomre ou d'un oligomre choisi dans l'ensemble constitu par les acrylates d'ther de nonylphnyle et de polyalkylneglycol, les mthacrylates d'ther de nonylphnyle et de polyalkylneglycol, et des mlanges de ceux-ci.

tous les pourcentages indiqus sont des pourcentages en poids par rapport au poids total de (A), (B) et
(C),

le revtement  l'tat durci ayant un indice de rfraction de 1,48 ou plus.

25 2. Revtement durcissable par rayonnement selon la revendication 1, dans lequel ledit oligomre (A) est caractris par la formule

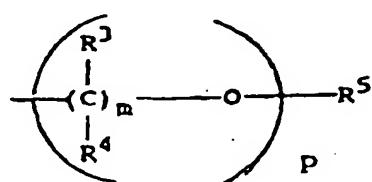


35 o

R¹ est un polymre hydrocarbon linaire ou ramifi ayant un poids molculaire de 500  4000, choisi dans l'ensemble constitu par un 1,2-polybutadine totalement ou partiellement hydrogn, un 1,2-polybutadine hydrogn jusqu' un indice d'iode de 9  21, un polyisobutylne totalement ou partiellement hydrogn, un polybutne-polyol et un diol dimre hydrogn ;40 R² est un alkylne linaire, ramifi ou cyclique, ayant de six  vingt atomes de carbone ; et

X et Q sont indpendamment :

45 (a) un radical rpondant  la formule :

50 o R⁵ est choisi dans l'ensemble constitu par les groupes acryliques, mthacryliques, vinyliques, allyliques, itaconiques, crotoniques, styrniques, acrylamides, norbornnyles, acetylniques, epoxy, mercapto et amino ; R³ et R⁴ sont indpendamment l'hydrogne ou un groupe mthyle, thyle ou propyle ; m est un nombre entier de 1  10 ; et p vaut zro ou un, ou

55 (b) un radical alkyle satur ayant de neuf  vingt atomes de carbone, avec la condition que ledit oligomre

dove posséder au moins un groupe terminal apte à réagir.

3. Revêtement durcissable par rayonnement selon la revendication 1 ou la revendication 2, dans lequel ledit monomère (B) est choisi dans l'ensemble constitué par l'acrylate d'hexyle, le méthacrylate d'hexyle, l'acrylate de 2-éthylhexyle, le méthacrylate de 2-éthylhexyle, l'acrylate d'octyle, le méthacrylate d'octyle, l'acrylate d'isoctyle, le méthacrylate d'isoctyle, l'acrylate de décyle, le méthacrylate de décyle, l'acrylate d'isodécyle, le méthacrylate d'isodécyle, l'acrylate de lauryle, le méthacrylate de lauryle, l'acrylate de stéaryl, le méthacrylate de stéaryl, les diacrylates de diols hydrocarbonés en C₁₄-C₁₅, les diméthacrylates de diols hydrocarbonés en C₁₄-C₁₅, l'acrylate de tridécyle, le méthacrylate de tridécyle, l'acrylate palmitique, le méthacrylate palmitique, le 1,10-décanedithiol, l'éther de divinyle et d'hexanediol, les esters mercapto-acétates, les esters mercaptopropionates, l'éther de vinyle et de lauryle, l'éther de vinyle et d'éthylhexyle, l'éther de vinyle et d'isodécyle, l'éther de divinyle et de butane-diol, l'éther de divinyle et de cyclohexanediméthanol, les éthers de divinyle et de diols en C₁₄-C₁₅, et des mélanges de ceux-ci.
4. Revêtement durcissable par rayonnement selon l'une quelconque des revendications 1 à 3, comprenant, en outre, de 1,0 % à 50,0 % en poids, par rapport au poids total de (A), (B) et (C), d'un agent de réticulation (D).
5. Revêtement durcissable par rayonnement selon la revendication 4, dans lequel ledit agent de réticulation (D) est l'éther de trivinyle et de triméthyloléthane.
6. Revêtement durcissable par rayonnement selon l'une quelconque des revendications 1 à 5, comprenant, en outre, de 0,0001 % à 3,0 % en poids, par rapport au poids total de (A), (B) et (C), d'un stabilisant (E).
7. Revêtement durcissable par rayonnement selon la revendication 6, dans lequel ledit stabilisant (E) est choisi dans l'ensemble constitué par la diéthylhydroxylamine, les phosphites organiques, les phénols empêchés, la trihexylamine, le 3-(3',5'-di-tert-butyl-4'-hydroxyphényl)propionate d'octadécyle, le bis(3,5-di-tert-butyl-4-hydroxy)hydrocinnamate de thiodiéthylène, un copolymère butylé de paracrésol-dicyclopentadiène, le tétrakis[méthylène-(3,5-di-tert-butyl-4-hydroxyhydrocinnamate]-méthane, le 3-aminopropyltriméthoxysilane, et des mélanges de ceux-ci.
8. Revêtement durcissable par rayonnement selon l'une quelconque des revendications 1 à 7, comprenant, en outre, de 0,1 % à 3,0 % en poids, par rapport au poids total de (A), (B) et (C), d'un silane organofonctionnel comme promoteur d'adhésion (F).
9. Revêtement durcissable par rayonnement selon l'une quelconque des revendications 1 à 8, dans lequel ledit photoinitiateur (C) est choisi dans l'ensemble constitué par l'hydroxyméthylphénylpropanone, l'hydroxycyclohexylphénylcétone, la diméthoxyphénylacétophénone, le 2-méthyl-1,4-(méthylthio)phényl-2-morpholino-propane-1, la 1-(4-isopropylphényl)-2-hydroxy-2-méthyl-propane-1-one, la 1-(4-dodécylphényl)-2-hydroxy-2-méthylpropane-1-one, la 4-(2-hydroxyéthoxy)phényl-2-(2-hydroxy-2-propyl)cétone, la diéthoxyacétophénone, la 2,2-di-sec-butoxyacétophénone, la diéthoxyphénylacétophénone, l'hexafluoroantimoniate de (4-octyloxypheényl)phényliodonium, l'hexafluoroantimoniate de (4-octyloxypheényl)diphenylsulfonium, l'hexafluoroantimoniate de (4-décyloxypheényl)phényliodonium, l'hexafluoroantimoniate de (4-octadécyloxypheényl)phényliodonium, et des mélanges de ceux-ci.
10. Revêtement durcissable par rayonnement selon l'une quelconque des revendications 1 à 9, comprenant, en outre, de 0,1 % à 10 % en poids, par rapport au poids total de (A), (B) et (C), d'un agent de transfert de chaîne (H) à fonction mercapto.
11. Revêtement durcissable par rayonnement selon l'une quelconque des revendications 1 à 10, qui est un revêtement primaire pour une fibre optique.
12. Revêtement durcissable par rayonnement selon la revendication 1, qui est un revêtement primaire pour une fibre optique, dans lequel les composants suivants sont présents :
 - (A) de 30 % à 70 % en poids d'un oligomère aliphatique d'uréthane et d'éther vinylique, ayant, comme squelette, un polymère de 1,2-polybutadiène hydrogéné ;
 - (B) de 10 % à 40 % en poids de 1,10-décanedithiol ;
 - (C) de 2 % à 10 % en poids d'hydroxyméthylphénylpropanone comme photo-initiateur ;
 - (D) de 1 % à 20 % en poids d'éther de trivinyle et de triméthyloléthane comme agent de réticulation ; et

(E) de 0,001 % à 0,1 % en poids de diéthylhydroxylamine comme stabilisant,

et dans lequel tous les pourcentages indiqués sont des pourcentages en poids se rapportant au poids total de (A), (B) et (C).

5 13. Revêtement durcissable par rayonnement selon la revendication 1, qui est un revêtement primaire pour une fibre optique, dans lequel les composants suivants sont présents :

10 (A) de 20 % à 80 % en poids d'un oligomère d'uréthane aliphatique à terminaison éther vinylique, ayant, comme squelette, un polymère de 1,2-butadiène hydrogéné ;
 (B) de 10 % à 80 % en poids d'éther de divinyle et d'hexanediol ; et
 (C) de 0,05 % à 5,0 % en poids d'hexafluoroantimonate de (4-octyloxyphényl)phényliodonium comme initiateur de durcissement cationique,

15 et dans lequel tous les pourcentages indiqués sont des pourcentages en poids se rapportant au poids total de (A), (B) et (C).

20 14. Fibre optique revêtue comprenant une fibre optique et une couche de revêtement primaire selon l'une quelconque des revendications 1 à 13.

25 15. Fibre optique revêtue selon la revendication 14, dans laquelle ladite couche de revêtement primaire a été durcie par rayonnement après son application sur ladite fibre de verre.

30 16. Fibre optique revêtue selon la revendication 14, dans laquelle ladite couche de revêtement primaire est formée à partir de matériaux ayant des groupes terminaux aptes à réagir, qui sont durcissables par rayonnement et qui sont durcis par rayonnement *in situ*.

35 17. Procédé de préparation d'une fibre optique revêtue, comprenant :

30 (i) l'application, sur une fibre optique, d'une couche de revêtement primaire comprenant une composition selon l'une quelconque des revendications 1 à 13 ; et
 (ii) l'application d'une couche de revêtement secondaire sur ladite couche de revêtement primaire ; et
 (iii) le durcissement par rayonnement *in situ* desdites couches de revêtements primaire et secondaire.

35 18. Procédé selon la revendication 17, dans lequel la couche de revêtement primaire comprend :

40 (A) de 45 % à 65 % en poids d'un oligomère d'acrylate d'uréthane aliphatique ayant, comme squelette, un polymère de 1,2-polybutadiène hydrogéné ;
 (B) de 10 % à 20 % en poids d'acrylate de lauryle ;
 (C) de 15 % à 30 % en poids d'acrylate d'éther de nonylphényle et de polypropylèneglycol ;
 (D) de 0,3 % à 1,0 % en poids de gamma-mercaptopropyltriméthoxysilane comme promoteur d'adhésion ; et
 (E) de 2,0 % à 7,0 % en poids d'hydroxycyclohexylphénylcétone comme photo-initiateur,

45 dans laquelle tous les pourcentages indiqués sont des pourcentages en poids se rapportant au poids total de (A), (B) et (C).

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